## AMÉNDMENT & RESPONSE UNDER 37 C.F.R. § 1.116 - EXPEDITED PROCEDURE

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Title: NON-TOXIC MUTANTS OF PATHOGENIC GRAM-NEGATIVE BACTERIA

Page 2 Dkt: 875.001US2



- 23. (Amended) A mutant endotoxin made according to the method of claim 22, wherein the mutant endotoxin was purified from the *htrB* mutant <u>pathogen</u> by phenol-water extraction or by protease digestion.
- 24. The mutant endotoxin according to claim 23, wherein the mutant endotoxin is conjugated to a carrier protein.
- 25. A mutant endotoxin made according to the method of claim 22.
- 26. The mutant endotoxin according to claim 25, wherein the mutant endotoxin is conjugated to a carrier protein.

29. (Amended) A method for producing endotoxin-specific antisera [for use in diagnostic assays], the method comprising

- immunizing an individual with a vaccine formulation comprising [as an active ingredient] an htrB mutant of a gram-negative bacterial pathogen, endotoxin isolated from the htrB mutant of the gram-negative bacterial pathogen, [and] or endotoxin [isolated] purified from the htrB mutant of the gram-negative bacterial pathogen wherein the endotoxin is conjugated to a carrier protein; and
- (b) collecting antibody produced from the immunized individual; wherein the htrB mutant lacks one or more secondary acyl chains of lipid A contained in [the] a wild type gram-negative bacterial pathogen resulting in substantially reduced toxicity when compared to lipid A of the wild type gram-negative bacterial pathogen.

Please add the following new claims 44 and 45:

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(New) The method of claim 22 wherein the gram-negative bacterial pathogen is of the genera Haemophilus, Neisseria, Moraxella, Campylobacter, Shigella or Pseudomonas.

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